

5                   **SYSTEM AND METHOD FOR DIAGNOSING AND MONITORING**  
                    **RESPIRATORY INSUFFICIENCY FOR AUTOMATED**  
                    **REMOTE PATIENT CARE**

**Cross-Reference to Related Application**

                    This patent application is a continuation of U.S. patent application, Serial  
No. 10/152,207, filed May 20, 2002, pending; which is a continuation of U.S.  
10   Patent No. 6,398,728, issued June 4, 2002, the disclosures of which are  
incorporated by reference.

**Field of the Invention**

                    The present invention relates in general to respiratory insufficiency  
diagnosis and analysis, and, in particular, to an automated collection and analysis  
15   patient care system and method for diagnosing and monitoring respiratory  
insufficiency and outcomes thereof throughout disease onset, progression,  
regression, and status quo.

**Background of the Invention**

                    Presently, respiratory insufficiency due to primary diseases of the lungs is  
20   one of the leading causes of acute and chronic illness in the world. Clinically,  
respiratory insufficiency involves either difficulty in ventilation or in  
oxygenation. The former is manifest by increases in the arterial partial pressure  
of carbon dioxide and the latter is manifest by decreases in arterial partial pressure  
of oxygen. For purposes of this invention, the term "respiratory insufficiency"  
25   will refer to ventilatory insufficiency and/or to problems in oxygenation due to  
diseases of the lung. Common causes of respiratory insufficiency include  
bronchitis, emphysema, pneumonia, pulmonary emboli, congestive heart failure,  
tumor infiltration of the lung and abnormalities of the interstitium of the lungs  
that may be infectious in origin, due to immunological abnormalities, or as a

result of exposure to environmental pathogens. The effects of respiratory insufficiency range from cough to impairment during physical exertion to a complete failure of lung function and respiratory arrest at any level of activity. Clinical manifestations of respiratory insufficiency include respiratory distress, such as shortness of breath and fatigue, cough, and reduced exercise capacity or tolerance.

Several factors make the early diagnosis and prevention of respiratory insufficiency, as well as the monitoring of the progression of respiratory insufficiency, relatively difficult. First, the onset of respiratory insufficiency is generally subtle and erratic. Often, the symptoms are ignored and the patient compensates by changing his or her daily activities. This situation is especially true in chronic lung disorders where the onset of symptoms can be very gradual. As a result, many respiratory insufficiency conditions or deteriorations in respiratory insufficiency remain undiagnosed until more serious problems arise seriously limiting the activities of daily living.

The susceptibility to suffer from respiratory insufficiency depends upon the patient's age, sex, physical condition, and other factors, such as smoking history, occupation, diabetes, co-existing heart disease, immunodepression, the presence or absence of cancer, surgical history, kidney function, and extent of pre-existing lung disease. No one factor is dispositive. Finally, annual or even monthly lung checkups, including chest X-rays or other lung tests, provide, at best, a "snapshot" of patient wellness and the incremental and subtle clinicophysiological changes which portend the onset or progression of respiratory insufficiency often go unnoticed, even with regular health care. Documentation of subtle improvements following therapy that can guide and refine further evaluation and therapy can be equally elusive.

Nevertheless, taking advantage of frequently and regularly measured physiological measures, such as recorded manually by a patient, via an external monitoring or therapeutic device, or via implantable device technologies, can provide a degree of detection and prevention heretofore unknown. For instance, patients already suffering from some form of treatable heart disease often receive

an implantable pulse generator (IPG), cardiovascular or heart failure monitor, therapeutic device, or similar external wearable device, with which rhythm and structural problems of the heart can be monitored and treated. These types of devices, although usually originally intended for use in treating some type of cardiac problem, can contain sufficient physiological data to allow accurate assessment of lung disorders. Such devices are useful for detecting physiological changes in patient conditions through the retrieval and analysis of telemetered signals stored in an on-board, volatile memory. Typically, these devices can store more than thirty minutes of per heartbeat and respiratory cycle data recorded on a per heartbeat, per respiration, binned average basis, or on a derived basis from, for example, atrial or ventricular electrical activity, minute ventilation, patient activity score, cardiac output score, arterial or mixed venous oxygen score, cardiopulmonary pressure measures, and the like. However, the proper analysis of retrieved telemetered signals requires detailed medical subspecialty knowledge, particularly by pulmonologists and cardiologists.

Alternatively, these telemetered signals can be remotely collected and analyzed using an automated patient care system. One such system is described in a related, commonly assigned U.S. Patent No. 6,312,378, issued November 6, 2001, the disclosure of which is incorporated herein by reference. A medical device adapted to be implanted in an individual patient records telemetered signals that are then retrieved on a regular, periodic basis using an interrogator or similar interfacing device. The telemetered signals are downloaded via an internetwork onto a network server on a regular, e.g., daily, basis and stored as sets of collected measures in a database along with other patient care records. The information is then analyzed in an automated fashion and feedback, which includes a patient status indicator, is provided to the patient.

While such an automated system can serve as a valuable tool in providing remote patient care, an approach to systematically correlating and analyzing the raw collected telemetered signals, as well as manually collected physiological measures, through applied pulmonary and cardiovascular medical knowledge to accurately diagnose the onset of a particular medical condition, such as

respiratory insufficiency, is needed, especially in patients with co-existing heart disease. One automated patient care system directed to a patient-specific monitoring function is described in U.S. Patent No. 5,113,869 ('869) to Nappholz et al. The '869 patent discloses an implantable, programmable

5 electrocardiography (ECG) patient monitoring device that senses and analyzes ECG signals to detect ECG and physiological signal characteristics predictive of malignant cardiac arrhythmias. The monitoring device can communicate a warning signal to an external device when arrhythmias are predicted. However, the Nappholz device is limited to detecting tachycardias. Unlike requirements for

10 automated respiratory insufficiency monitoring, the Nappholz device focuses on rudimentary ECG signals indicative of malignant cardiac tachycardias, an already well established technique that can be readily used with on-board signal detection techniques. Also, the Nappholz device is patient specific only and is unable to automatically take into consideration a broader patient or peer group history for

15 reference to detect and consider the progression or improvement of lung disease. Moreover, the Nappholz device has a limited capability to automatically self-reference multiple data points in time and cannot detect disease regression even in the individual patient. Also, the Nappholz device must be implanted and cannot function as an external monitor. Finally, the Nappholz device is incapable of

20 tracking the cardiovascular and cardiopulmonary consequences of any rhythm disorder.

Consequently, there is a need for a systematic approach to detecting trends in regularly collected physiological data indicative of the onset, progression, regression, or status quo of respiratory insufficiency diagnosed and monitored

25 using an automated, remote patient care system. The physiological data could be telemetered signals data recorded either by an external or an implantable medical device or, alternatively, individual measures collected through manual means. Preferably, such an approach would be capable of diagnosing both acute and chronic respiratory insufficiency conditions, as well as the symptoms of other

30 lung disorders. In addition, findings from individual, peer group, and general

population patient care records could be integrated into continuous, on-going monitoring and analysis.

### **Summary of the Invention**

The present invention provides a system and method for diagnosing and monitoring the onset, progression, regression, and status quo of respiratory insufficiency using an automated collection and analysis patient care system. Measures of patient cardiopulmonary information are either recorded by an external or implantable medical device, such as an IPG, cardiovascular or heart failure monitor, or respiratory diagnostic or therapeutic device, or manually through conventional patient-operable means. The measures are collected on a regular, periodic basis for storage in a database along with other patient care records. Derived measures are developed from the stored measures. Select stored and derived measures are analyzed and changes in patient condition are logged. The logged changes are compared to quantified indicator thresholds to detect findings of respiratory distress or reduced exercise capacity indicative of the principal pathophysiological manifestations of respiratory insufficiency: elevated partial pressure of arterial carbon dioxide and reduced partial pressure of arterial oxygen.

An embodiment of the present invention is an automated system and method for diagnosing and monitoring respiratory insufficiency and outcomes thereof. A plurality of monitoring sets is retrieved from a database. Each of the monitoring sets include recorded measures relating to patient information recorded on a substantially continuous basis. A patient status change is determined by comparing at least one recorded measure from each of the monitoring sets to at least one other recorded measure. Both recorded measures relate to the same type of patient information. Each patient status change is tested against an indicator threshold corresponding to the same type of patient information as the recorded measures that were compared. The indicator threshold corresponds to a quantifiable physiological measure of a pathophysiology indicative of respiratory insufficiency.

A further embodiment is an automated collection and analysis patient care system and method for diagnosing and monitoring respiratory insufficiency and outcomes thereof. A plurality of monitoring sets is retrieved from a database. Each monitoring set includes recorded measures that each relate to patient  
5 information and include either medical device measures or derived measures calculable therefrom. The medical device measures are recorded on a substantially continuous basis. A set of indicator thresholds is defined. Each indicator threshold corresponds to a quantifiable physiological measure of a pathophysiology indicative of respiratory insufficiency and relates to the same  
10 type of patient information as at least one of the recorded measures. A respiratory insufficiency finding is diagnosed. A change in patient status is determined by comparing at least one recorded measure to at least one other recorded measure with both recorded measures relating to the same type of patient information. Each patient status change is compared to the indicator threshold corresponding to  
15 the same type of patient information as the recorded measures that were compared.

A further embodiment is an automated patient care system and method for diagnosing and monitoring respiratory insufficiency and outcomes thereof. Recorded measures organized into a monitoring set for an individual patient are  
20 stored into a database. Each recorded measure is recorded on a substantially continuous basis and relates to at least one aspect of monitoring reduced exercise capacity and/or respiratory distress. A plurality of the monitoring sets is periodically retrieved from the database. At least one measure related to respiratory insufficiency onset, progression, regression, and status quo is  
25 evaluated. A patient status change is determined by comparing at least one recorded measure from each of the monitoring sets to at least one other recorded measure with both recorded measures relating to the same type of patient information. Each patient status change is tested against an indicator threshold corresponding to the same type of patient information as the recorded measures  
30 that were compared. The indicator threshold corresponds to a quantifiable

physiological measure of a pathophysiology indicative of reduced exercise capacity and/or respiratory distress.

5       The present invention provides a capability to detect and track subtle trends and incremental changes in recorded patient cardiopulmonary information for diagnosing and monitoring respiratory insufficiency. When coupled with an enrollment in a remote patient monitoring service having the capability to remotely and continuously collect and analyze external or implantable medical device measures, respiratory insufficiency detection, prevention and tracking regression from therapeutic maneuvers become feasible.

10       Still other embodiments of the present invention will become readily apparent to those skilled in the art from the following detailed description, wherein is described embodiments of the invention by way of illustrating the best mode contemplated for carrying out the invention. As will be realized, the invention is capable of other and different embodiments and its several details are  
15       capable of modifications in various obvious respects, all without departing from the spirit and the scope of the present invention. Accordingly, the drawings and detailed description are to be regarded as illustrative in nature and not as restrictive.

#### **Brief Description of the Drawings**

20       FIGURE 1 is a block diagram showing an automated collection and analysis patient care system for diagnosing and monitoring respiratory insufficiency and outcomes thereof in accordance with the present invention;

25       FIGURE 2 is a database schema showing, by way of example, the organization of a device and derived measures set record for care of patients with respiratory insufficiency stored as part of a patient care record in the database of the system of FIGURE 1;

30       FIGURE 3 is a database schema showing, by way of example, the organization of a quality of life and symptom measures set record for care of patients with respiratory insufficiency stored as part of a patient care record in the database of the system of FIGURE 1;

FIGURE 4 is a database schema showing, by way of example, the organization of a combined measures set record for care of patients with respiratory insufficiency stored as part of a patient care record in the database of the system of FIGURE 1;

5           FIGURE 5 is a block diagram showing the software modules of the server system of the system of FIGURE 1;

FIGURE 6 is a record view showing, by way of example, a set of partial patient care records for care of patients with respiratory insufficiency stored in the database of the system of FIGURE 1;

10           FIGURE 7 is a Venn diagram showing, by way of example, peer group overlap between the partial patient care records of FIGURE 6;

FIGURES 8A-8B are flow diagrams showing a method for diagnosing and monitoring respiratory insufficiency and outcomes thereof using an automated collection and analysis patient care system in accordance with the present  
15   invention;

FIGURE 9 is a flow diagram showing the routine for retrieving reference baseline sets for use in the method of FIGURES 8A-8B;

FIGURE 10 is a flow diagram showing the routine for retrieving monitoring sets for use in the method of FIGURES 8A-8B;

20           FIGURES 11A-11F are flow diagrams showing the routine for testing threshold limits for use in the method of FIGURES 8A-8B;

FIGURE 12 is a flow diagram showing the routine for evaluating the onset, progression, regression, and status quo of respiratory insufficiency for use in the method of FIGURES 8A-8B;

25           FIGURES 13A-13C are flow diagrams showing the routine for determining an onset of respiratory insufficiency for use in the routine of FIGURE 12;

FIGURES 14A-14C are flow diagrams showing the routine for determining progression or worsening of respiratory insufficiency for use in the  
30   routine of FIGURE 12;



FIGURES 15A-15C are flow diagrams showing the routine for determining regression or improving of respiratory insufficiency for use in the routine of FIGURE 12; and

FIGURE 16 is a flow diagram showing the routine for determining  
5 threshold stickiness (“hysteresis”) for use in the method of FIGURE 12.

### **Detailed Description**

FIGURE 1 is a block diagram showing an automated collection and analysis patient care system 10 for diagnosing and monitoring respiratory insufficiency in accordance with the present invention. An exemplary automated  
10 collection and analysis patient care system suitable for use with the present invention is disclosed in the related, commonly assigned U.S. Patent No. 6,312,378, issued November 6, 2001, the disclosure of which is incorporated herein by reference. Preferably, an individual patient 11 is a recipient of an implantable medical device 12, such as, by way of example, an IPG,  
15 cardiovascular, heart failure monitor, pulmonary monitor, or therapeutic device, with a set of leads extending into his or her heart and electrodes implanted throughout the cardiopulmonary system. Alternatively, an external monitoring or therapeutic medical device 26, a subcutaneous monitor or device inserted into other organs, a cutaneous monitor, or even a manual physiological measurement  
20 device, such as an respiratory monitor, electrocardiogram or heart rate monitor, could be used. The implantable medical device 12 and external medical device 26 include circuitry for recording into a short-term, volatile memory telemetered signals stored for later retrieval, which become part of a set of device and derived measures, such as described below, by way of example, with reference to  
25 FIGURE 2. Exemplary implantable medical devices suitable for use in the present invention include the Discovery line of pacemakers, manufactured by Guidant Corporation, Indianapolis, Indiana, and the Gem line of ICDs, manufactured by Medtronic Corporation, Minneapolis, Minnesota.

The telemetered signals stored in the implantable medical device 12 are  
30 preferably retrieved upon the completion of an initial observation period and subsequently thereafter on a continuous, periodic (daily) basis, such as described

in the related, commonly assigned U.S. Patent No. 6,221,011, issued April 24, 2001, the disclosure of which is incorporated herein by reference. A programmer 14, personal computer 18, or similar device for communicating with an implantable medical device 12 can be used to retrieve the telemetered signals. A magnetized reed switch (not shown) within the implantable medical device 12 closes in response to the placement of a wand 13 over the site of the implantable medical device 12. The programmer 14 sends programming or interrogating instructions to and retrieves stored telemetered signals from the implantable medical device 12 via RF signals exchanged through the wand 13. Similar communication means are used for accessing the external medical device 26. Once downloaded, the telemetered signals are sent via an internetwork 15, such as the Internet, to a server system 16 which periodically receives and stores the telemetered signals as device measures in patient care records 23 in a database 17, as further described below, by way of example, with reference to FIGURES 2 and 3. An exemplary programmer 14 suitable for use in the present invention is the Model 2901 Programmer Recorder Monitor, manufactured by Guidant Corporation, Indianapolis, Indiana.

The patient 11 is remotely monitored by the server system 16 via the internetwork 15 through the periodic receipt of the retrieved device measures from the implantable medical device 12 or external medical device 26. The patient care records 23 in the database 17 are organized into two identified sets of device measures: an optional reference baseline 26 recorded during an initial observation period and monitoring sets 27 recorded subsequently thereafter. The device measures sets are periodically analyzed and compared by the server system 16 to indicator thresholds corresponding to quantifiable physiological measures of a pathophysiology indicative of respiratory insufficiency, as further described below with reference to FIGURE 5. As necessary, feedback is provided to the patient 11. By way of example, the feedback includes an electronic mail message automatically sent by the server system 16 over the internetwork 15 to a personal computer 18 (PC) situated for local access by the patient 11. Alternatively, the feedback can be sent through a telephone interface device 19 as an automated

voice mail message to a telephone 21 or as an automated facsimile message to a facsimile machine 22, both also situated for local access by the patient 11.

Moreover, simultaneous notifications can also be delivered to the patient's physician, hospital, or emergency medical services provider 29 using similar

5 feedback means to deliver the information.

The server system 10 can consist of either a single computer system or a cooperatively networked or clustered set of computer systems. Each computer system is a general purpose, programmed digital computing device consisting of a central processing unit (CPU), random access memory (RAM), non-volatile  
10 secondary storage, such as a hard drive or CD ROM drive, network interfaces, and peripheral devices, including user interfacing means, such as a keyboard and display. Program code, including software programs, and data are loaded into the RAM for execution and processing by the CPU and results are generated for display, output, transmittal, or storage, as is known in the art.

15 The database 17 stores patient care records 23 for each individual patient to whom remote patient care is being provided. Each patient care record 23 contains normal patient identification and treatment profile information, as well as medical history, medications taken, height and weight, and other pertinent data (not shown). The patient care records 23 consist primarily of two sets of data:  
20 device and derived measures (D&DM) sets 24a, 24b and quality of life (QOL) sets 25a, 25b, the organization of which are further described below with respect to FIGURES 2 and 3, respectively. The device and derived measures sets 24a, 24b and quality of life and symptom measures sets 25a, 25b can be further logically categorized into two potentially overlapping sets. The reference  
25 baseline 26 is a special set of device and derived reference measures sets 24a and quality of life and symptom measures sets 25a recorded and determined during an initial observation period. Monitoring sets 27 are device and derived measures sets 24b and quality of life and symptom measures sets 25b recorded and determined thereafter on a regular, continuous basis. Other forms of database  
30 organization are feasible.

The implantable medical device 12 and, in a more limited fashion, the external medical device 26, record patient information for care of patients with respiratory insufficiency on a regular basis. The recorded patient information is downloaded and stored in the database 17 as part of a patient care record 23.

- 5 Further patient information can be derived from recorded data, as is known in the art. FIGURE 2 is a database schema showing, by way of example, the organization of a device and derived measures set record 40 for patient care stored as part of a patient care record in the database 17 of the system of FIGURE 1. Each record 40 stores patient information which includes a snapshot of
- 10 telemetered signals data which were recorded by the implantable medical device 12 or the external medical device 26, for instance, on per heartbeat, binned average or derived bases; measures derived from the recorded device measures; and manually collected information, such as obtained through a patient medical history interview or questionnaire. The following non-exclusive information can
- 15 be recorded for a patient: atrial electrical activity 41, ventricular electrical activity 42, PR interval or AV interval 43, QRS measures 44, ST-T wave measures 45, QT interval 46, body temperature 47, patient activity score 48, posture 49, cardiovascular pressures 50, pulmonary artery systolic pressure measure 51, pulmonary artery diastolic pressure measure 52, respiratory rate 53, ventilatory
- 20 tidal volume 54, minute ventilation 55, transthoracic impedance 56, cardiac output 57, systemic blood pressure 58, patient geographic location (altitude) 59, mixed venous oxygen score 60, arterial oxygen score 61, arterial carbon dioxide score 62, acidity (pH) level 63, potassium [K+] level 64, sodium [Na+] level 65, glucose level 66, blood urea nitrogen (BUN) and creatinine 67, hematocrit 68,
- 25 hormonal levels 69, lung injury chemical tests 70, cardiac injury chemical tests 71, myocardial blood flow 72, central nervous system (CNS) injury chemical tests 73, central nervous system blood flow 74, interventions made by the implantable medical device or external medical device 75, and the relative success of any interventions made 76. In addition, the implantable medical device or external
- 30 medical device communicates device-specific information, including battery status, general device status and program settings 77 and the time of day 78 for

the various recorded measures. Other types of collected, recorded, combined, or derived measures are possible, as is known in the art.

The device and derived measures sets 24a, 24b (shown in FIGURE 1), along with quality of life and symptom measures sets 25a, 25b, as further  
5 described below with reference to FIGURE 3, are continuously and periodically received by the server system 16 as part of the on-going patient care monitoring and analysis function. These regularly collected data sets are collectively categorized as the monitoring sets 27 (shown in FIGURE 1). In addition, select device and derived measures sets 24a and quality of life and symptom measures  
10 sets 25a can be designated as a reference baseline 26 at the outset of patient care to improve the accuracy and meaningfulness of the serial monitoring sets 27. Select patient information is collected, recorded, and derived during an initial period of observation or patient care, such as described in the related, commonly assigned U.S. Patent No. 6,221,011, issued April 24, 2001, the disclosure of  
15 which is incorporated herein by reference.

As an adjunct to remote patient care through the monitoring of measured physiological data via the implantable medical device 12 or external medical device 26, quality of life and symptom measures sets 25a can also be stored in the database 17 as part of the reference baseline 26, if used, and the monitoring sets  
20 27. A quality of life measure is a semi-quantitative self-assessment of an individual patient's physical and emotional well being and a record of symptoms, such as provided by the Duke Activities Status Indicator. These scoring systems can be provided for use by the patient 11 on the personal computer 18 (shown in FIGURE 1) to record his or her quality of life scores for both initial and periodic  
25 download to the server system 16. FIGURE 3 is a database schema showing, by way of example, the organization of a quality of life record 80 for use in the database 17. The following information is recorded for a patient: overall health wellness 81, psychological state 82, activities of daily living 83, work status 84, geographic location 85, family status 86, shortness of breath 87, cough 88, sputum  
30 production 89, sputum color 90, energy level 91, exercise tolerance 92, chest

discomfort 93, and time of day 94, and other quality of life and symptom measures as would be known to one skilled in the art.

The patient may also add non-device quantitative measures, such as the six-minute walk distance, as complementary data to the device and derived  
5 measures sets 24a, 24b and the symptoms during the six-minute walk to quality of life and symptom measures sets 25a, 25b.

Other types of quality of life and symptom measures are possible, such as those indicated by responses to the Minnesota Living with Heart Failure Questionnaire described in E. Braunwald, ed., "Heart Disease—A Textbook of  
10 Cardiovascular Medicine," pp. 452-454, W.B. Saunders Co. (1997), the disclosure of which is incorporated herein by reference. Similarly, functional classifications based on the relationship between symptoms and the amount of effort required to provoke them can serve as quality of life and symptom measures, such as the New York Heart Association (NYHA) classifications I, II, III and IV, adapted for use  
15 for lung disease rather than heart disease, also described in *Ibid*.

On a periodic basis, the patient information stored in the database 17 is analyzed and compared to pre-determined cutoff levels, which, when exceeded, can provide etiological indications of respiratory insufficiency symptoms. FIGURE 4 is a database schema showing, by way of example, the organization of  
20 a combined measures set record 95 for use in the database 17. Each record 95 stores patient information obtained or derived from the device and derived measures sets 24a, 24b and quality of life and symptom measures sets 25a, 25b as maintained in the reference baseline 26, if used, and the monitoring sets 27. The combined measures set 95 represents those measures most (but not exhaustively  
25 or exclusively) relevant to a pathophysiology indicative of respiratory insufficiency and are determined as further described below with reference to FIGURES 8A-8B. The following information is stored for a patient: heart rate 96, heart rhythm (e.g., normal sinus vs. atrial fibrillation) 97, pacing modality 98, pulmonary artery systolic pressure measure 99, pulmonary artery diastolic  
30 pressure measure 100, cardiac output score 101, arterial oxygen score 102, mixed venous oxygen score 103, respiratory rate 104, tidal volume 105, transthoracic

impedance 106, arterial carbon dioxide score 107, right ventricular peak systolic pressure 108, pulmonary artery end diastolic pressure 109, patient activity score 110, posture 111, exercise tolerance quality of life and symptom measures 112, respiratory distress quality of life and symptom measures 113, cough 114, sputum  
5 production 115, any interventions made to treat respiratory insufficiency 116, including treatment by medical device, via drug infusion administered by the patient or by a medical device, surgery, and any other form of medical intervention as is known in the art, the relative success of any such interventions made 117, and time of day 118. Other types of comparison measures regarding  
10 respiratory insufficiency are possible as is known in the art. In the described embodiment, each combined measures set 95 is sequentially retrieved from the database 17 and processed. Alternatively, each combined measures set 95 could be stored within a dynamic data structure maintained transitorily in the random access memory of the server system 16 during the analysis and comparison  
15 operations.

FIGURE 5 is a block diagram showing the software modules of the server system 16 of the system 10 of FIGURE 1. Each module is a computer program written as source code in a conventional programming language, such as the C or Java programming languages, and is presented for execution by the CPU of the  
20 server system 16 as object or byte code, as is known in the art. The various implementations of the source code and object and byte codes can be held on a computer-readable storage medium or embodied on a transmission medium in a carrier wave. The server system 16 includes three primary software modules, database module 125, diagnostic module 126, and feedback module 128, which  
25 perform integrated functions as follows.

First, the database module 125 organizes the individual patient care records 23 stored in the database 17 (shown in FIGURE 1) and efficiently stores and accesses the reference baseline 26, monitoring sets 27, and patient care data maintained in those records. Any type of database organization could be utilized,  
30 including a flat file system, hierarchical database, relational database, or

distributed database, such as provided by database vendors, such as Oracle Corporation, Redwood Shores, California.

Next, the diagnostic module 126 makes findings of respiratory insufficiency based on the comparison and analysis of the data measures from the reference baseline 26 and monitoring sets 27. The diagnostic module includes  
5 three modules: comparison module 130, analysis module 131, and quality of life module 132. The comparison module 130 compares recorded and derived measures retrieved from the reference baseline 26, if used, and monitoring sets 27 to indicator thresholds 129. The database 17 stores individual patient care records  
10 23 for patients suffering from various health disorders and diseases for which they are receiving remote patient care. For purposes of comparison and analysis by the comparison module 130, these records can be categorized into peer groups containing the records for those patients suffering from similar disorders, as well as being viewed in reference to the overall patient population. The definition of  
15 the peer group can be progressively refined as the overall patient population grows. To illustrate, FIGURE 6 is a record view showing, by way of example, a set of partial patient care records for care of patients with respiratory insufficiency stored in the database 17 for three patients, *Patient 1*, *Patient 2*, and *Patient 3*. For each patient, three sets of peer measures, *X*, *Y*, and *Z*, are shown. Each of the  
20 measures, *X*, *Y*, and *Z*, could be either collected or derived measures from the reference baseline 26, if used, and monitoring sets 27.

The same measures are organized into time-based sets with *Set 0* representing sibling measures made at a reference time  $t=0$ . Similarly, *Set n-2*, *Set n-1* and *Set n* each represent sibling measures made at later reference times  
25  $t=n-2$ ,  $t=n-1$  and  $t=n$ , respectively. Thus, for a given patient, such as *Patient 1*, serial peer measures, such as peer measure  $X_0$  through  $X_n$ , represent the same type of patient information monitored over time. The combined peer measures for all patients can be categorized into a health disorder- or disease-matched peer group. The definition of disease-matched peer group is a progressive definition, refined  
30 over time as the number of monitored patients grows. Measures representing different types of patient information, such as measures  $X_0$ ,  $Y_0$ , and  $Z_0$ , are sibling



measures. These are measures which are also measured over time, but which might have medically significant meaning when compared to each other within a set for an individual patient.

The comparison module 130 performs two basic forms of comparisons.

- 5 First, individual measures for a given patient can be compared to other individual measures for that same patient (self-referencing). These comparisons might be peer-to-peer measures, that is, measures relating to a one specific type of patient information, projected over time, for instance,  $X_n, X_{n-1}, X_{n-2}, \dots X_0$ , or sibling-to-sibling measures, that is, measures relating to multiple types of patient  
10 information measured during the same time period, for a single snapshot, for instance,  $X_n, Y_n$ , and  $Z_n$ , or projected over time, for instance,  $X_n, Y_n, Z_n, X_{n-1}, Y_{n-1}, Z_{n-1}, X_{n-2}, Y_{n-2}, Z_{n-2}, \dots X_0, Y_0, Z_0$ . Second, individual measures for a given patient can be compared to other individual measures for a group of other patients sharing the same disorder- or disease-specific characteristics (peer group  
15 referencing) or to the patient population in general (population referencing). Again, these comparisons might be peer-to-peer measures projected over time, for instance,  $X_n, X_n', X_n'', X_{n-1}, X_{n-1}', X_{n-1}'', X_{n-2}, X_{n-2}', X_{n-2}'' \dots X_0, X_0', X_0''$ , or comparing the individual patient's measures to an average from the group. Similarly, these comparisons might be sibling-to-sibling measures for single  
20 snapshots, for instance,  $X_n, X_n', X_n'', Y_n, Y_n', Y_n'',$  and  $Z_n, Z_n', Z_n''$ , or projected over time, for instance,  $X_n, X_n', X_n'', Y_n, Y_n', Y_n'', Z_n, Z_n', Z_n'', X_{n-1}, X_{n-1}', X_{n-1}'', Y_{n-1}, Y_{n-1}', Y_{n-1}'', Z_{n-1}, Z_{n-1}', Z_{n-1}'', X_{n-2}, X_{n-2}', X_{n-2}'', Y_{n-2}, Y_{n-2}', Y_{n-2}'', Z_{n-2}, Z_{n-2}', Z_{n-2}'' \dots X_0, X_0', X_0'', Y_0, Y_0', Y_0'',$  and  $Z_0, Z_0', Z_0''$ . Other forms of comparisons are feasible, including multiple disease diagnoses for diseases exhibiting similar abnormalities  
25 in physiological measures that result from a second disease but manifest in different combinations or onset in different temporal sequences.

FIGURE 7 is a Venn diagram showing, by way of example, peer group overlap between the partial patient care records 23 of FIGURE 1. Each patient care record 23 includes characteristics data 350, 351, 352, including personal  
30 traits, demographics, medical history, and related personal data, for patients 1, 2 and 3, respectively. For example, the characteristics data 350 for patient 1 might

include personal traits which include gender and age, such as male and an age between 40-45; a demographic of resident of New York City; and a medical history consisting of chronic bronchitis, recurrent pneumonia, a history of an inferior myocardial infarction and diabetes. Similarly, the characteristics data 351 for patient 2 might include identical personal traits, thereby resulting in partial overlap 353 of characteristics data 350 and 351. Similar characteristics overlap 354, 355, 356 can exist between each respective patient. The overall patient population 357 would include the universe of all characteristics data. As the monitoring population grows, the number of patients with personal traits matching those of the monitored patient will grow, increasing the value of peer group referencing. Large peer groups, well matched across all monitored measures, will result in a well known natural history of disease and will allow for more accurate prediction of the clinical course of the patient being monitored. If the population of patients is relatively small, only some traits 356 will be uniformly present in any particular peer group. Eventually, peer groups, for instance, composed of 100 or more patients each, would evolve under conditions in which there would be complete overlap of substantially all salient data, thereby forming a powerful core reference group for any new patient being monitored.

Referring back to FIGURE 5, the analysis module 131 analyzes the results from the comparison module 130, which are stored as a combined measures set 95 (not shown), to a set of indicator thresholds 129, as further described below with reference to FIGURES 8A-8B. Similarly, the quality of life module 132 compares quality of life and symptom measures 25a, 25b from the reference baseline 26 and monitoring sets 27, the results of which are incorporated into the comparisons performed by the analysis module 131, in part, to either refute or support the findings based on physiological "hard" data. Finally, the feedback module 128 provides automated feedback to the individual patient based, in part, on the patient status indicator 127 generated by the diagnostic module 126. As described above, the feedback could be by electronic mail or by automated voice mail or facsimile. The feedback can also include normalized voice feedback, such as described in the related, commonly assigned U.S. Patent No. 6,203,495, issued

March 20, 2001, the disclosure of which is incorporated herein by reference. In addition, the feedback module 128 determines whether any changes to interventive measures are appropriate based on threshold stickiness ("hysteresis") 133, as further described below with reference to FIGURE 16. The threshold stickiness 133 can prevent fickleness in the diagnostic routines resulting from transient, non-trending and non-significant fluctuations in the various collected and derived measures in favor of more certainty in diagnosis. However, in the case of some of the parameters being followed, such as activity and pulmonary artery systolic and diastolic pressures, abrupt spikes in these measures can be indicative of coughing and therefore helpful in indicating the onset of pulmonary insufficiency. In a further embodiment of the present invention, the feedback module 128 includes a patient query engine 134 that enables the individual patient 11 to interactively query the server system 16 regarding the diagnosis, therapeutic maneuvers, and treatment regimen. Conversely, the patient query engines 134, found in interactive expert systems for diagnosing medical conditions, can interactively query the patient. Using the personal computer 18 (shown in FIGURE 1), the patient can have an interactive dialogue with the automated server system 16, as well as human experts as necessary, to self assess his or her medical condition. Such expert systems are well known in the art, an example of which is the MYCIN expert system developed at Stanford University and described in Buchanan, B. & Shortlife, E., "RULE-BASED EXPERT SYSTEMS. The MYCIN Experiments of the Stanford Heuristic Programming Project," Addison-Wesley (1984). The various forms of feedback described above help to increase the accuracy and specificity of the reporting of the quality of life and symptomatic measures.

FIGURES 8A-8B are flow diagrams showing a method for diagnosing and monitoring respiratory insufficiency and outcomes thereof 135 using an automated collection and analysis patient care system 10 in accordance with the present invention. First, the indicator thresholds 129 (shown in FIGURE 5) are set (block 136) by defining a quantifiable physiological measure of a pathophysiology indicative of respiratory insufficiency and relating to the each

type of patient information in the combined device and derived measures set 95 (shown in FIGURE 4). The actual values of each indicator threshold can be finite cutoff values, weighted values, or statistical ranges, as discussed below with reference to FIGURES 11A-11F. Next, the reference baseline 26 (block 137) and  
5 monitoring sets 27 (block 138) are retrieved from the database 17, as further described below with reference to FIGURES 9 and 10, respectively. Each measure in the combined device and derived measures set 95 is tested against the threshold limits defined for each indicator threshold 129 (block 139), as further described below with reference to FIGURES 11A-11F. The potential onset,  
10 progression, regression, or status quo of respiratory insufficiency is then evaluated (block 140) based upon the findings of the threshold limits tests (block 139), as further described below with reference to FIGURES 13A-13C, 14A-14C, 15A-15C.

In a further embodiment, multiple near-simultaneous disorders are  
15 considered in addition to primary respiratory insufficiency. *Primary respiratory insufficiency* is defined as the onset or progression of respiratory insufficiency without obvious inciting cause. *Secondary respiratory insufficiency* is defined as the onset or progression of respiratory insufficiency (in a patient with or without pre-existing respiratory insufficiency) from another disease process, such as  
20 congestive heart failure, coronary insufficiency, atrial fibrillation, and so forth. Other health disorders and diseases can potentially share the same forms of symptomatology as respiratory insufficiency, such as congestive heart failure, myocardial ischemia, pneumonia, exacerbation of chronic bronchitis, renal failure, sleep-apnea, stroke, anemia, atrial fibrillation, other cardiac arrhythmias,  
25 and so forth. If more than one abnormality is present, the relative sequence and magnitude of onset of abnormalities in the monitored measures becomes most important in sorting and prioritizing disease diagnosis and treatment.

Thus, if other disorders or diseases are being cross-referenced and diagnosed (block 141), their status is determined (block 142). In the described  
30 embodiment, the operations of ordering and prioritizing multiple near-simultaneous disorders (box 151) by the testing of threshold limits and analysis in

a manner similar to congestive heart failure as described above, preferably in parallel to the present determination, is described in the related, commonly assigned U.S. Patent No. 6,440,066, entitled "Automated Collection And Analysis Patient Care System And Method For Ordering And Prioritizing Multiple Health Disorders To Identify An Index Disorder," issued August 27, 2002, the disclosure of which is incorporated herein by reference. If respiratory insufficiency is due to an obvious inciting cause, i.e., secondary respiratory insufficiency, (block 143), an appropriate treatment regimen for respiratory insufficiency as exacerbated by other disorders is adopted that includes treatment of secondary disorders, e.g., congestive heart failure, myocardial ischemia, atrial fibrillation, and so forth (block 144) and a suitable patient status indicator 127 for respiratory insufficiency is provided (block 146) to the patient. Suitable devices and approaches to diagnosing and treating congestive heart failure, myocardial infarction, and atrial fibrillation are described in related, commonly assigned U.S. Patent No. 6,336,903, entitled "Automated Collection And Analysis Patient Care System And Method For Diagnosing And Monitoring Congestive Heart Failure And Outcomes Thereof," issued January 8, 2002; U.S. Patent No. 6,368,284, entitled "Automated Collection And Analysis Patient Care System And Method For Diagnosing And Monitoring Myocardial Ischemia And Outcomes Thereof," issued April 9, 2002; and U.S. Patent No. 6,411,840, entitled "Automated Collection And Analysis Patient Care System And Method For Diagnosing And Monitoring The Outcomes Of Atrial Fibrillation," June 15, 2002, the disclosures of which are incorporated herein by reference.

Otherwise, if primary respiratory insufficiency is indicated (block 143), a primary treatment regimen is followed (block 145). A patient status indicator 127 for respiratory insufficiency is provided (block 146) to the patient regarding physical well-being, disease prognosis, including any determinations of disease onset, progression, regression, or status quo, and other pertinent medical and general information of potential interest to the patient.

Finally, in a further embodiment, if the patient submits a query to the server system 16 (block 147), the patient query is interactively processed by the

patient query engine (block 148). Similarly, if the server elects to query the patient (block 149), the server query is interactively processed by the server query engine (block 150). The method then terminates if no further patient or server queries are submitted.

5           FIGURE 9 is a flow diagram showing the routine for retrieving reference baseline sets 137 for use in the method of FIGURES 8A-8B. The purpose of this routine is to retrieve the appropriate reference baseline sets 26, if used, from the database 17 based on the types of comparisons being performed. First, if the comparisons are self referencing with respect to the measures stored in the individual patient care record 23 (block 152), the reference device and derived measures set 24a and reference quality of life and symptom measures set 25a, if used, are retrieved for the individual patient from the database 17 (block 153). Next, if the comparisons are peer group referencing with respect to measures stored in the patient care records 23 for a health disorder- or disease-specific peer group (block 154), the reference device and derived measures set 24a and reference quality of life and symptom measures set 25a, if used, are retrieved from each patient care record 23 for the peer group from the database 17 (block 155). Data for each measure (e.g., minimum, maximum, averaged, standard deviation (SD), and trending data) from the reference baseline 26 for the peer group is then calculated (block 156). Finally, if the comparisons are population referencing with respect to measures stored in the patient care records 23 for the overall patient population (block 157), the reference device and derived measures set 24a and reference quality of life and symptom measures set 25a, if used, are retrieved from each patient care record 23 from the database 17 (block 158). Minimum, maximum, averaged, standard deviation, and trending data and other numerical processes using the data, as is known in the art, for each measure from the reference baseline 26 for the peer group is then calculated (block 159). The routine then returns.

30           FIGURE 10 is a flow diagram showing the routine for retrieving monitoring sets 138 for use in the method of FIGURES 8A-8B. The purpose of this routine is to retrieve the appropriate monitoring sets 27 from the database 17

based on the types of comparisons being performed. First, if the comparisons are self referencing with respect to the measures stored in the individual patient care record 23 (block 160), the device and derived measures set 24b and quality of life and symptom measures set 25b, if used, are retrieved for the individual patient  
5 from the database 17 (block 161). Next, if the comparisons are peer group referencing with respect to measures stored in the patient care records 23 for a health disorder- or disease-specific peer group (block 162), the device and derived measures set 24b and quality of life and symptom measures set 25b, if used, are retrieved from each patient care record 23 for the peer group from the database 17  
10 (block 163). Data for each measure (e.g., minimum, maximum, averaged, standard deviation, and trending data) from the monitoring sets 27 for the peer group is then calculated (block 164). Finally, if the comparisons are population referencing with respect to measures stored in the patient care records 23 for the overall patient population (block 165), the device and derived measures set 24b  
15 and quality of life and symptom measures set 25b, if used, are retrieved from each patient care record 23 from the database 17 (block 166). Minimum, maximum, averaged, standard deviation, and trending data and other numerical processes using the data, as is known in the art, for each measure from the monitoring sets 27 for the peer group is then calculated (block 167). The routine then returns.

20        FIGURES 11A-11F are flow diagrams showing the routine for testing threshold limits 139 for use in the method of FIGURE 8A and 8B. The purpose of this routine is to analyze, compare, and log any differences between the observed, objective measures stored in the reference baseline 26, if used, and the monitoring sets 27 to the indicator thresholds 129. Briefly, the routine consists of  
25 tests pertaining to each of the indicators relevant to diagnosing and monitoring respiratory insufficiency. The threshold tests focus primarily on: (1) changes to and rates of change for the indicators themselves, as stored in the combined device and derived measures set 95 (shown in FIGURE 4) or similar data structure; and (2) violations of absolute threshold limits which trigger an alert.  
30 The timing and degree of change may vary with each measure and with the natural fluctuations noted in that measure during the reference baseline period. In

addition, the timing and degree of change might also vary with the individual and the natural history of a measure for that patient.

One suitable approach to performing the threshold tests uses a standard statistical linear regression technique using a least squares error fit. The least

5 squares error fit can be calculated as follows:

$$y = \beta_0 + \beta_1 x \quad (1)$$

$$\beta = \frac{SS_{xy}}{SS_{xx}} \quad (2)$$

$$SS_{xy} = \sum_{i=1}^n x_i y_i - \frac{\left( \sum_{i=1}^n x_i \right) \left( \sum_{i=1}^n y_i \right)}{n} \quad (3)$$

$$SS_{xx} = \sum_{i=1}^n x_i^2 - \frac{\left( \sum_{i=1}^n x_i \right)^2}{n} \quad (4)$$

10 where  $n$  is the total number of measures,  $x_i$  is the time of day for measure  $i$ , and  $y_i$  is the value of measure  $i$ ,  $\beta_1$  is the slope, and  $\beta_0$  is the  $y$ -intercept of the least squares error line. A positive slope  $\beta_1$  indicates an increasing trend, a negative slope  $\beta_1$  indicates a decreasing trend, and no slope indicates no change in patient condition for that particular measure. A predicted measure value can be  
15 calculated and compared to the appropriate indicator threshold 129 for determining whether the particular measure has either exceeded an acceptable threshold rate of change or the absolute threshold limit.

For any given patient, three basic types of comparisons between individual measures stored in the monitoring sets 27 are possible: self referencing, peer  
20 group, and general population, as explained above with reference to FIGURE 6. In addition, each of these comparisons can include comparisons to individual measures stored in the pertinent reference baselines 24.

The indicator thresholds 129 for detecting a trend indicating progression into a state of respiratory insufficiency or a state of imminent or likely respiratory  
25 insufficiency, for example, over a one week time period, can be as follows:



- (1) Respiratory rate (block 170): If the respiratory rate has increased over 1.0 SD from the mean respiratory rate in the reference baseline 26 (block 171), the increased respiratory rate and time span over which it occurs are logged in the combined measures set 95 (block 172).
- (2) Heart rate (block 173): If the heart rate has increased over 1.0 SD from the mean heart rate in the reference baseline 26 (block 174), the increased heart rate and time span over which it occurs are logged in the combined measures set 95 (block 175).
- (3) Transthoracic impedance (block 176): If the transthoracic impedance has increased over 1.0 SD from the mean transthoracic impedance in the reference baseline 26 (block 177), the increased transthoracic impedance and time span are logged in the combined measures set 95 (block 178).
- (4) The ventilatory tidal volume (block 179): If the tidal volume has increased over 1.0 SD from the tidal volume score in the reference baseline 26 (block 180), the increased tidal volume score and time span are logged in the combined measures set 95 (block 181).
- (5) Arterial oxygen score (block 182): If the arterial oxygen score has decreased over 1.0 SD from the arterial oxygen score in the reference baseline 26 (block 183), the decreased arterial oxygen score and time span are logged in the combined measures set 95 (block 184).
- (6) Arterial carbon dioxide score (block 185): If the arterial carbon dioxide score has decreased over 1.0 SD from the arterial carbon dioxide score in the reference baseline 26 (block 186), the decreased arterial carbon dioxide score and time span are logged in the combined measures set 95 (block 187).
- (7) Patient activity score (block 188): If the mean patient activity score has decreased over 1.0 SD from the mean patient activity score in the reference baseline 26 (block 189), the decreased patient activity

score and time span are logged in the combined measures set 95 (block 190).

5 (8) Temperature (block 191): If the patient temperature score has increased over 1.0 SD from the mean patient temperature score in the reference baseline 26 (block 192), the increased patient temperature score and the time span are logged in the combined measures set 95 (block 193).

10 (9) Spikes in patient activity (block 194): If short-lived spikes in the patient activity score occur over time periods under 5 minutes compared to the reference baseline 26 (block 195), the spike in patient activity score and time span are logged in the combined measures set 95 (block 196).

15 (10) Spikes in pulmonary arterial pressure (PAP) (block 197): If short-lived spikes in the pulmonary arterial pressure score occur over time periods under 5 minutes compared to the reference baseline 26 (block 198), the spike in the pulmonary arterial pressure score and time span are logged in the combined measures set 95 (block 199). In the described embodiment, the mean arterial pressure on any spike in the arterial pressure tracing could be utilized.

20 (11) Exercise tolerance quality of life (QOL) measures (block 200): If the exercise tolerance QOL has decreased over 1.0 SD from the mean exercise tolerance in the reference baseline 26 (block 201), the decrease in exercise tolerance and the time span over which it occurs are logged in the combined measures set 95 (block 202).

25 (12) Respiratory distress quality of life (QOL) measures (block 203): If the respiratory distress QOL measure has deteriorated by more than 1.0 SD from the mean respiratory distress QOL measure in the reference baseline 26 (block 204), the increase in respiratory distress and the time span over which it occurs are logged in the combined measures set 95 (block 205).

30

- (13) Spikes in right ventricular (RV) pressure (block 206): If short-lived spikes in the right ventricular pressure occur over time periods under 5 minutes compared to the reference baseline 26 (block 207), the spike in the right ventricular pressure and time span are logged in the combined measures set 95 (block 208).
- (14) Spikes in transthoracic impedance (TTZ) (block 209): If short-lived spikes in the transthoracic impedance occur over time periods under 5 minutes compared to the reference baseline 26 (block 210), the spike in the transthoracic impedance and time span are logged in the combined measures set 95 (block 211).
- (15) Atrial fibrillation (block 212): The presence or absence of atrial fibrillation (AF) is determined and, if present (block 213), atrial fibrillation is logged (block 214).
- (16) Rhythm changes (block 215): The type and sequence of rhythm changes is significant and is determined (block 216) based on the timing of the relevant rhythm measure, such as sinus rhythm. For instance, a finding that a rhythm change to atrial fibrillation precipitated respiratory measures changes can indicate therapy directions against atrial fibrillation rather than the primary development of respiratory insufficiency. Thus, if there are rhythm changes (block 217), the sequence of the rhythm changes and time span are logged (block 211).

Note also that an inversion of the indicator thresholds 129 defined above could similarly be used for detecting a trend in disease regression. One skilled in the art would recognize that these measures would vary based on whether or not they were recorded during rest or during activity and that the measured activity score can be used to indicate the degree of patient rest or activity. The patient activity score can be determined via an implantable motion detector, for example, as described in U.S. Patent No. 4,428,378, issued January 31, 1984, to Anderson et al., the disclosure of which is incorporated herein by reference.

The indicator thresholds 129 for detecting a trend towards a state of respiratory insufficiency can also be used to declare, *a priori*, respiratory insufficiency present, regardless of pre-existing trend data when certain limits are established, such as:

- 5           (1)    An absolute limit of arterial oxygen (block 182) less than 85 mm Hg is an *a priori* definition of respiratory insufficiency from decreased oxygenation.
- (2)    An absolute limit of arterial carbon dioxide (block 185) falling  
10           below 25 mm Hg (in the absence of marked exercise) or greater  
            than 50 mm Hg are both *a priori* definitions of respiratory  
            insufficiency as indicated by hyperventilation and hypoventilation,  
            respectively.

FIGURE 12 is a flow diagram showing the routine for evaluating the onset, progression, regression and status quo of respiratory insufficiency 140 for  
15   use in the method of FIGURE 8A and 8B. The purpose of this routine is to  
evaluate the presence of sufficient indicia to warrant a diagnosis of the onset,  
progression, regression, and status quo of respiratory insufficiency. Quality of  
life and symptom measures 25a, 25b can be included in the evaluation (block 230)  
by determining whether any of the individual quality of life and symptom  
20   measures 25a, 25b have changed relative to the previously collected quality of life  
and symptom measures from the monitoring sets 27 and the reference baseline 26,  
if used. For example, an increase in the shortness of breath measure 87 and  
exercise tolerance measure 92 would corroborate a finding of respiratory  
insufficiency. Similarly, a transition from NYHA Class II to NYHA Class III  
25   would indicate a deterioration or, conversely, a transition from NYHA Class III to  
NYHA Class II status would indicate improvement or progress when adapting the  
NYHA classifications for their parallel in lung disorders. Incorporating the  
quality of life and symptom measures 25a, 25b into the evaluation can help, in  
part, to refute or support findings based on physiological data. Next, a  
30   determination as to whether any changes to interventive measures are appropriate

based on threshold stickiness (“hysteresis”) is made (block 231), as further described below with reference to FIGURE 16.

The routine returns upon either the determination of a finding or elimination of all factors as follows. If a finding of respiratory insufficiency was not previously diagnosed (block 232), a determination of disease onset is made (block 233), as further described below with reference to FIGURES 13A-13C. Otherwise, if respiratory insufficiency was previously diagnosed (block 232), a further determination of either disease progression or worsening (block 234) or regression or improving (block 235) is made, as further described below with reference to FIGURES 14A-14C and 15A-15C, respectively. If, upon evaluation, neither disease onset (block 233), worsening (block 234) or improving (block 235) is indicated, a finding of status quo is appropriate (block 236) and noted (block 237). Otherwise, respiratory insufficiency and the related outcomes are actively managed (block 238) through the administration of, non-exclusively, antibiotic and antiviral therapies, bronchodilator therapies, oxygen therapies, antiinflammation therapies, electrical therapies, mechanical therapies, and other therapies as are known in the art. The management of respiratory insufficiency is described, by way of example, in A.S. Fauci et al. (Eds.), “Harrison’s Principles of Internal Medicine,” pp. 1407-1491, McGraw-Hill, 14<sup>th</sup> Ed. (1997), the disclosure of which is incorporated herein by reference. The routine then returns.

FIGURES 13A-13C are flow diagrams showing the routine for determining an onset of respiratory insufficiency 232 for use in the routine of FIGURE 12. Respiratory insufficiency is possible based on two general symptom categories: reduced exercise capacity (block 244) and respiratory distress (block 256). An effort is made to diagnose respiratory insufficiency manifesting primarily as resulting in reduced exercise capacity (block 244) and/or increased respiratory distress (block 256). Reduced exercise capacity and respiratory distress can generally serve as markers of low systemic arterial oxygenation. The clinical aspects of respiratory insufficiency are described, by way of example, in A.S. Fauci et al. (Eds.), “Harrison’s Principles of Internal Medicine,” pp. 1410-

1419, McGraw-Hill, 14<sup>th</sup> Ed. (1997), the disclosure of which is incorporated herein by reference.

As primary pulmonary disease considerations, multiple individual indications (blocks 240-243, 245-253) should be present for the two principal findings of respiratory insufficiency related reduced exercise capacity (block 244), or respiratory insufficiency related respiratory distress (block 256), to be indicated, both for disease onset or progression. The presence of primary key findings alone can be sufficient to indicate an onset of respiratory insufficiency and secondary key findings serve to corroborate disease onset. Note the presence of any abnormality can trigger an analysis for the presence or absence of secondary disease processes, such as the presence of atrial fibrillation or congestive heart failure. Secondary disease considerations can be evaluated using the same indications (see, e.g., blocks 141-144 of FIGURES 8A-8B), but with adjusted indicator thresholds 129 (shown in FIGURE 5) triggered at a change of 0.5 SD, for example, instead of 1.0 SD.

In the described embodiment, the reduced exercise capacity and respiratory distress findings (blocks 244, 251) can be established by consolidating the individual indications (blocks 240-243, 245-253) in several ways. First, in a preferred embodiment, each individual indication (blocks 240-243, 245-253) is assigned a scaled index value correlating with the relative severity of the indication. For example, decreased cardiac output (block 240) could be measured on a scale from '1' to '5' wherein a score of '1' indicates no change in cardiac output from the reference point, a score of '2' indicates a change exceeding 0.5 SD, a score of '3' indicates a change exceeding 1.0 SD, a score of '4' indicates a change exceeding 2.0 SD, and a score of '5' indicates a change exceeding 3.0 SD. The index value for each of the individual indications (blocks 240-243, 245-253) can then either be aggregated or averaged with a result exceeding the aggregate or average maximum indicating an appropriate respiratory insufficiency finding.

Preferably, all scores are weighted depending upon the assignments made from the measures in the reference baseline 26. For instance, arterial partial

pressure of oxygen 102 could be weighted more importantly than respiratory rate 104 if the respiratory rate in the reference baseline 26 is particularly high at the outset, making the detection of further disease progression from increases in respiratory rate, less sensitive. In the described embodiment, arterial partial pressure of oxygen 102 receives the most weight in determining a reduced exercise capacity finding whereas arterial partial pressure of carbon dioxide 107 receives the most weight in determining a respiratory distress or dyspnea finding.

Alternatively, a simple binary decision tree can be utilized wherein each of the individual indications (blocks 240-243, 245-253) is either present or is not present. All or a majority of the individual indications (blocks 240-243, 245-253) should be present for the relevant respiratory insufficiency finding to be affirmed.

Other forms of consolidating the individual indications (blocks 240-243, 245-253) are feasible.

FIGURES 14A-14C are flow diagrams showing the routine for determining a progression or worsening of respiratory insufficiency 234 for use in the routine of FIGURE 12. The primary difference between the determinations of disease onset, as described with reference to FIGURES 13A-13C, and disease progression is the evaluation of changes indicated in the same factors present in a disease onset finding. Thus, a revised respiratory insufficiency finding is possible based on the same two general symptom categories: reduced exercise capacity (block 274) and respiratory distress (block 286). The same factors which need be indicated to warrant a diagnosis of respiratory insufficiency onset are evaluated to determine disease progression.

Similarly, FIGURES 15A-15C are flow diagrams showing the routine for determining a regression or improving of respiratory distress 235 for use in the routine of FIGURE 12. The same factors as described above with reference to FIGURES 13A-13C and 14A-14C, trending in opposite directions from disease onset or progression, are evaluated to determine disease regression. As primary cardiac disease considerations, multiple individual indications (blocks 300-303, 305-313) should be present for the two principal findings of respiratory insufficiency related reduced exercise capacity (block 304), or respiratory

insufficiency related respiratory distress (block 316), to indicate disease regression.

FIGURE 16 is a flow diagram showing the routine for determining threshold stickiness (“hysteresis”) 231 for use in the method of FIGURE 12.

5 Stickiness, also known as hysteresis, is a medical practice doctrine whereby a diagnosis or therapy will not be changed based upon small or temporary changes in a patient reading, even though those changes might temporarily move into a new zone of concern. For example, if a patient measure can vary along a scale of ‘1’ to ‘10’ with ‘10’ being worse, a transient reading of ‘6,’ standing alone, on a  
10 patient who has consistently indicated a reading of ‘5’ for weeks will not warrant a change in diagnosis without a definitive prolonged deterioration first being indicated. Stickiness dictates that small or temporary changes require more diagnostic certainty, as confirmed by the persistence of the changes, than large changes would require for any of the monitored (device) measures. Stickiness  
15 also makes reversal of important diagnostic decisions, particularly those regarding life-threatening disorders, more difficult than reversal of diagnoses of modest import. As an example, automatic external defibrillators (AEDs) manufactured by Heartstream, a subsidiary of Agilent Technologies, Seattle, Washington, monitor heart rhythms and provide interventive shock treatment for the diagnosis  
20 of ventricular fibrillation. Once diagnosis of ventricular fibrillation and a decision to shock the patient has been made, a pattern of no ventricular fibrillation must be indicated for a relatively prolonged period before the AED changes to a “no-shock” decision. As implemented in this AED example, stickiness mandates certainty before a decision to shock is disregarded. In practice, stickiness also  
25 dictates that acute deteriorations in disease state are treated aggressively while chronic, more slowly progressing disease states are treated in a more tempered fashion. However, in the case of some of the parameters being followed, such as activity and pulmonary artery systolic pressure, abrupt spikes in these measures can be indicative of coughing and therefore helpful in indicating the onset of a  
30 disorder that might lead to pulmonary insufficiency.



Thus, if the patient status indicates a status quo (block 330), no changes in treatment or diagnosis are indicated and the routine returns. Otherwise, if the patient status indicates a change away from status quo (block 330), the relative quantum of change and the length of time over which the change has occurred is determinative. If the change of approximately 0.5 SD has occurred over the course of about one month (block 331), a gradually deteriorating condition exists (block 332) and a very tempered diagnostic, and if appropriate, treatment program is undertaken. If the change of approximately 1.0 SD has occurred over the course of about one week (block 333), a more rapidly deteriorating condition exists (block 334) and a slightly more aggressive diagnostic, and if appropriate, treatment program is undertaken. If the change of approximately 2.0 SD has occurred over the course of about one day (block 335), an urgently deteriorating condition exists (block 336) and a moderately aggressive diagnostic, and if appropriate, treatment program is undertaken. If the change of approximately 3.0 SD has occurred over the course of about one hour (block 337), an emergency condition exists (block 338) and an immediate diagnostic, and if appropriate, treatment program is undertaken as is practical. Finally, if the change and duration fall outside the aforementioned ranges (blocks 331-338), an exceptional condition exists (block 339) and the changes are reviewed manually, if necessary. The routine then returns. These threshold limits and time ranges may then be adapted depending upon patient history and peer-group guidelines.

The present invention provides several benefits. One benefit is improved predictive accuracy from the outset of patient care when a reference baseline is incorporated into the automated diagnosis. Another benefit is an expanded knowledge base created by expanding the methodologies applied to a single patient to include patient peer groups and the overall patient population. Collaterally, the information maintained in the database could also be utilized for the development of further predictive techniques and for medical research purposes. Yet a further benefit is the ability to hone and improve the predictive techniques employed through a continual reassessment of patient therapy outcomes and morbidity rates.

Other benefits include an automated, expert system approach to the cross-referral, consideration, and potential finding or elimination of other diseases and health disorders with similar or related etiological indicators and for those other disorders that may have an impact on respiratory insufficiency. Although disease  
5 specific markers will prove very useful in discriminating the underlying cause of symptoms, many diseases, other than respiratory insufficiency, will alter some of the same physiological measures indicative of respiratory insufficiency. Consequently, an important aspect of considering the potential impact of other disorders will be, not only the monitoring of disease specific markers, but the  
10 sequencing of change and the temporal evolution of more general physiological measures, for example respiratory rate, pulmonary artery diastolic pressure, and cardiac output, to reflect disease onset, progression or regression in more than one type of disease process, especially congestive heart failure from whatever cause.

While the invention has been particularly shown and described as  
15 referenced to the embodiments thereof, those skilled in the art will understand that the foregoing and other changes in form and detail may be made therein without departing from the spirit and scope of the invention.